EXHIBIT 1

ANDREW GOBEA:

RECEIVED 160 cc CORD BLOOD (40 CPD)
RBC mono 30%
Poly 70%

added 15 ml 1xPBS to bring to incubation volume of 75ml.
Added 1.5 ml antibody (12.8)
incubated 25ml.
Primed CellPro "cprate".

Spun cells. Resuspended in 1x PBS to a final volume of 300cc in bag.

Ran through column.

Unadsorbed portion → spun down. and consolidated in 1xPBS for incubation.

75ml for incubation (added HSA)
1.5ml antibody (12.8) 25min. Spun down following incubation. Resd to Vol. of 300cc in bag
Ran through 2nd column.

Stem cell portions from Runs 1 & 2 were combined (after counts done and samples removed for staining)

Total cells 2.8x10^6 for transduction

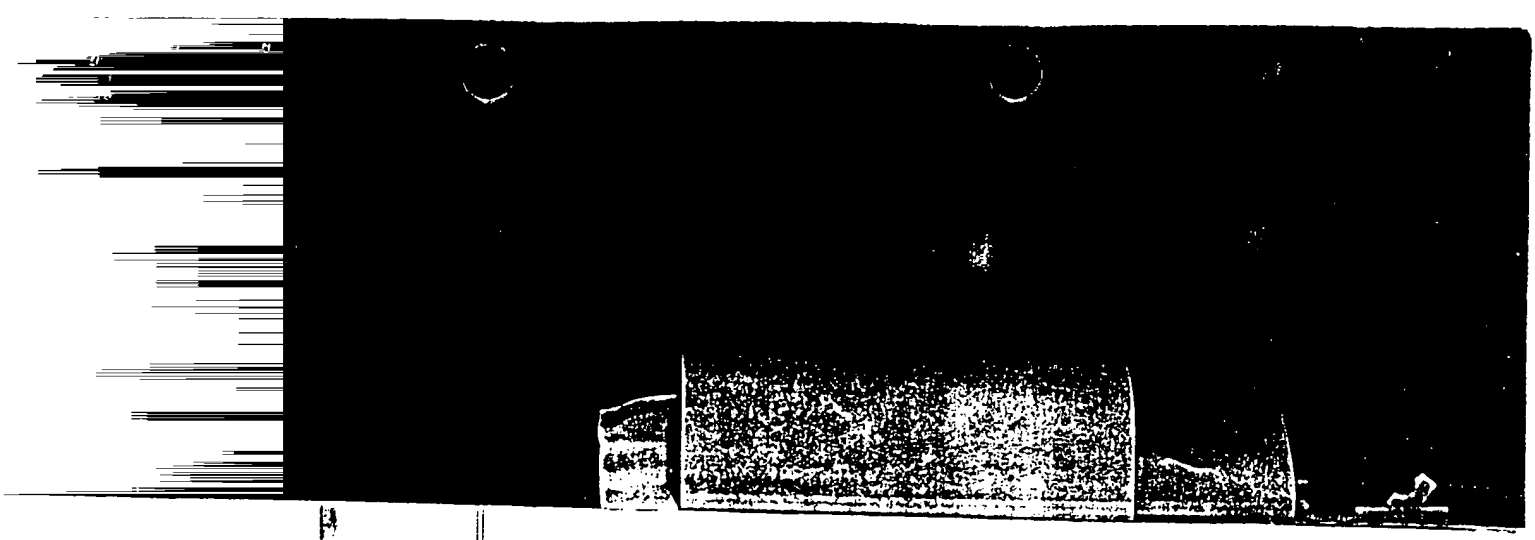
BBMM + 3

IL3 SANDO
Wash
IL6 SANDO
Wash
SCF AMGE
Wash

final concer
will be dil
Cells are
concentrat

BBMM: FB
BS

2.8x10^6 cells
Put in



BBMM + 316ISCF (for 500ml of media).

IL3 SANDOZ # Y0230892 stock at 150ug/ml
want final: 20ng/ml x 2 .. 20ug 133ul add

IL6 SANDOZ # Y0150392 stock at 150ug/ml
want final: 50ng/ml x 2 .. 50ug 333ul add

SCF AMGEN # 1509F2 stock at 1.5mg/ml = 1500ug/ml
want final: 100ng/ml x 2 .. 100ug 67ul add

final concentrations are doubled since the media will be diluted 1:2 w/ viral supernatant. Cells are therefore incubated with the correct concentrations.

BBMM: FBS Genuni LOT# A9603H
BSA #115

2.8x10⁶ cells want final: 5x10⁴ cells.
Put in 2 T-75 30ml each: 15ml BBMS 051193
15ml LAS VEGAS LOT# 53
+ protamine sulfate 240ul of 1:10 diluted stock

Cord Blood cells pre processing:

CFUs SCT 143

	Plate #	Sample	# Cells	# ul/ml media
-G418	1ab	Gabea UCB	5×10^4	50
+G418	2ab	↓	↓	50
-G418	3ab		1×10^5	100
+G418	4ab		↓	100

Start:

absorbed fraction:

absorbed fraction

CFUs Post transduction: SCT 144

	plate #	# CELLS	# ul
-G418	1ab	500	7
↓	2ab	1000	14
	3ab	2000	28
+G418	4ab	500	7
↓	5ab	1000	14
	6ab	2000	28

(yields) absorbed

↑ count:

$\bar{x} = 34$

$\times 2 \times 10^4 = 6.8 \times 10^5$ Cfu/ml

$\times 5.5$ ml $= 3.7 \times 10^6$ C

Refused on 5/15/93

No transduced stem cells

G418			
&	7ab	1000	20
+	8ab	2000	40

absorbed

Start:

	Pre	Post ab
5×10^8 C	0.71%	0.22%

*34+ = 3.6×10^6 C = 1.1×10^6 C

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absorbed fraction #1:

	FL1 FL2 gate	FL1 FSC gate
2×10^6 C	31.94%	20.81%
*34+ =	0.64×10^6 C	0.42×10^6 C

absorbed fraction #2:

0.8×10^6 C	2.46%	5.80%
*34+c =	0.02×10^6 C	0.05×10^6 C

(yields)

absorbed #1:

Pre & FL1/FL2 gate	Pre & FL1/FSC gate
$\frac{0.64 \times 10^6 c}{3.6 \times 10^6 C}$ = 17.8%	$\frac{0.42 \times 10^6 c}{3.6 \times 10^6 C}$ = 11.7%
Post ab & FL1/FL2 gate	**Post ab & FL1/FSC gate**
$\frac{0.64 \times 10^6 c}{1.1 \times 10^6 C}$ = 58.2%	$\frac{0.42 \times 10^6 c}{1.1 \times 10^6 C}$ = 38.2%

absorbed #2: Pre & FL1/FL2 | Pre & FL1/FSC

[illegible] from [illegible]

ZACHARY RIGGINS: 5/14/93

RECD 200cc COLD BLOOD

PRE: mono 109 ✓ poly 109

218 x 50 = 10.9 x 10⁶ c/ml
x 200ml = 2.2 x 10⁹ c start

Added 3 vials (4.5ml) 12.8 ab
inc. 25 min.

Spun down. Rspd. in 1xPBS to 300ml in bag.

Ran through column:

Spun down unadsorbed fraction for 2nd ab incubation.

Spun stem cell fraction to Rspd in smaller volume for count.

COUNTS:

unadsorbed

mono 107 ✓ poly 102

169 x 50 x 10³
= 8.5 x 10⁶ c/ml x 225ml
= 1.9 x 10⁹ c

stem

mono 172 ✓ poly 16

188 x 2 x 10⁴
= 3.8 x 10⁶ c/ml x 5.5ml
= 20.7 x 10⁶ c

incubated unadsorbed fraction w/ 4.5 mL
12.8 ab. for 25 min.
Spun down.
Rxn in 300 mL in bag (w/ 1x PBS)
Ran through 2nd column.

Counts:

unadsorbed		stem	
mono	poly	mono	poly
30	33	58	4

$63 \times 53 \times 10^3$
3.15×10^6 c/mL x 600 mL
$= 1.9 \times 10^9$ c

$62 \times 2 \times 10^4$
$= 1.2 \times 10^6$ c/mL
$\times 5$ mL $= 6 \times 10^6$ c

⇓

percolled/ficolled
free ⇒ LOT 12)

combined stem cell fractions

26×10^6 c for transduction
want final [] = 5×10^4 c/mL ... 520 mL total
÷ 2 = 260 mL supe
260 mL media

13 flasks 40 mL/flask 20 mL supe
20 mL media (B365)
+ 300 uL protamine sulfate
(12)

LASN supe 539 (bottles 18/19)

CFUs:

5C7145

PRE

Plate #	Example	# Cells	# ml
	(-G418)		
1ab	PRE trans	5×10^4	5
2ab	↓ (+G418)	↓	5

BBMM + 3161SCF:

1L3 Sandoz # yo230292

1L6 Sandoz # yo150392

SCF AMGEN # 150952

Took sample to micro for sterility ✓
each day of transduction pd. 0
Stat Gram stain done (negative)
before cells were given to baby.

5/15/93 4pm 2nd transduction:
Spun cells down from each flask
Rspd in fresh media & LASN supe
added Protamine sulfate

5/16/93 3rd transduction 330pm.
Repeated above.

5/17/93 cells washed 4x
3x in 1xPBS + PI's
last wash in RPMI (no PI's)

count: 160×10^6 c $\bar{x} = 15 \times 10 \times 10^4 = 15 \times 10^6$ c
$\times 40\,ml = 160 \times 10^6$ c

Put in 5cc into 10cc syringe

Refused on 5/17/93 (UCSF)

Post trans. CFU's: sq146

sample plate #		# cells	# µl
DG418	1ab	500	4
↓	2ab	1000	8
↓	3ab	2000	16
+G418	4ab	500	4
↓	5ab	1000	8
↓	6ab	2000	16
DG418	7ab	1000	24
+G418	8ab	1000	24

µl

5
5

y ✓
strike)
baby
tion
h flask
SN supe

30pm.